Ocular Therapeutix™ to Report Fourth Quarter and Full Year 2015 Financial Results

March 2, 2016

BEDFORD, Mass.--(BUSINESS WIRE)--Mar. 2, 2016-- Ocular Therapeutix, Inc. (NASDAQ:OCUL), a biopharmaceutical company focused on the development and commercialization of innovative therapies for diseases and conditions of the eye, today announced that it will report financial results for the fourth quarter and full year ended December 31, 2015, on Thursday, March 10, 2016. Following distribution of the earnings release via wire services, the Ocular management team will host a live conference call and webcast at 8:00 a.m. Eastern Time to review the Company's financial results and provide a business update.

The live webcast can be accessed by visiting the investor section of Ocular's website at investors.ocutx.com. Please connect to the Company's website at least 15 minutes prior to the live webcast to ensure adequate time for any software download that may be needed to access the webcast. Alternatively, please call 844-464-3934 (U.S.) or 765-507-2620 (International) to listen to the conference call. The conference ID number for the live call will be 60154048. An archive of the webcast will be available until March 24, 2016 on the company's website.

About Ocular Therapeutix, Inc.

Ocular Therapeutix, Inc. (NASDAQ: OCUL) is a biopharmaceutical company focused on the development and commercialization of innovative therapies for diseases and conditions of the eye using its proprietary hydrogel platform technology. Ocular Therapeutix's lead product candidate, DEXTENZA™, is in Phase 3 clinical development for post-surgical ocular inflammation and pain and allergic conjunctivitis, and in Phase 2 clinical development for inflammatory dry eye disease. An NDA for the post-operative ocular pain indication has been accepted by the FDA with a PDUFA date of July 24, 2016. A third Phase 3 clinical trial is being conducted for post-operative ocular inflammation and pain. For glaucoma and ocular hypertension, OTX-TP (sustained release travoprost) intracanalicular depot, has completed a Phase 2b clinical trial, and plans to commence the first of two planned Phase 3 clinical trials in the third quarter of 2016. Ocular Therapeutix is also evaluating sustained-release injectable anti-VEGF drug depots for back-of-the-eye diseases. Ocular Therapeutix's first product, ReSure® Sealant, is FDA-approved to seal corneal incisions following cataract surgery.

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Source: Ocular Therapeutix, Inc.

Investors
Ocular Therapeutix, Inc.
Brad Smith
Chief Financial Officer
bsmith@ocutx.com
or
Burns McClellan on behalf of Ocular Therapeutix
Steve Klass, 212-213-0006
sklass@burnsmc.com
or
Media
Ocular Therapeutix, Inc.
Scott Corning
Vice President of Sales and Marketing
scorning@ocutx.com